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Protein for blocking platelet adhesion

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Protein for blocking platelet adhesion

Summary of the invention

5

A naturally occurring protein isolated from the saliva of the medicinal leech *Hirudo medicinalis* is described which strongly binds to collagen thus acting as an inhibitor of natural platelet adhesion to collagen. The protein has a molecular weight of about 12 000 Dalton, an acidic isoelectric point and contains six
10 cysteins. The protein was sequenced and the gene was cloned from a *H. medicinalis* cDNA-library. Procedures for producing such polypeptide by recombinant techniques are disclosed. The recombinant and the natural occurring proteins are potent inhibitors of collagen-dependent platelet adhesion and therefore useful for the therapeutic treatment of various conditions related to hard disease and
15 diseases of the circulation system. Furthermore the protein is useful for coating natural or artificial collagen surfaces in order to render them nonadhesive for cells and prevent the activation of cells.

Field of invention

20

In haemostasis or thrombosis, platelets adhere to the cell-extracellular matrix of an injured vessel and cover the surface of the damaged area. Preventing this important initial step in the pathogenesis of thrombosis and arterial occlusion should be of therapeutic benefit in the effort of to prevent thrombotic diseases.
25 Collagen is considered to be the most thrombogenic surface component and has been shown to be a strong stimulant for platelet adhesion, aggregation and the release of their granules leading to the recruitment of (Ruggeri, Z.M. et al.; Seminars in Hematology, 1994, 31, 229-39) additional platelets to this area to form aggregates or a thrombus. The initial contact of the platelets to the vessel
30 surface is mediated by collagen bound von Willebrand Factor (vWF) and a specific vWF receptor on platelets, the glycoprotein Ib-V-IX complex. In addition ADP, epinephrine and circulating clotting factors drive the further activation process of platelets while simultaneously an increase in thrombin activity

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contributes to the formation of the cross-linked fibrin clot. Platelet-platelet aggregation supports this process and is mainly driven by fibrinogen as a mediator that bridges cells through the glycoprotein IIb/IIIa receptor.

- 5 This normal physiological response is quite critical in the course of the pathological process where platelets adhere to collagenes exposed in sclerotic lesions (Van der Rest M. et al.; FASEB Journal, 1991, 5, 2814-23) and start to build-up occlusions. Depending on the location and extent of the occlusion severe complications such as myocardial infarction, stroke, inflammation or pulmonary
10 embolism may be the severe outcome of this process.

- As a direct acting antithrombotic agent heparin which blocks the thrombin activity, thus preventing the formation of fibrin rich thrombi, is the currently most well known drug used in anti-thrombotic interventions. Heparin is widely used in
15 indications such as: unstable angina and acute myocardial infarction. However despite the wide use several severe short comings such as intravenous application, requirement for anti-thrombin-III as a cofactor, reduced affinity for clot-bound thrombin, its inactivation by several plasma proteins, the occasional induction of thrombocytopenia and its biological heterogeneity remain
20 unresolved. As a consequence the results of using heparin in the clinical setting have not been overwhelming so far.

- Recent development of low molecular weight heparin has contributed a version for subcutaneous application, however the therapeutic benefit over the standard
25 heparin has been modest. Unfortunately the same applies to the other directly acting antithrombins such as Hirudin, Hirulog and Warfarin. It turned out, that one of the major problems seems to be related to the increased production of thrombin under antithrombotic treatment (Rao, A.K et al., Circulation, 1996, 94, 389-2395).

- 30 Other recent strategies have therefore been focussed to the process of prothrombin activation which is driven by Factor Xa. The major challenge is the design of appropriate inhibitors directed to this factor. In summary one would

therefore argue that the full therapeutic potential of this type of intervention has not yet been realized.

Another pannel of therapeutics is represented by the thrombolytic regimens and has been focussed on the development of staphylokinase, streptokinase, 5 urokinase type Plasminogen Activator, tissue type Plasminogen Activator and anisoylated-plasminogen-streptokinase activator complex. The differences in time necessary to inducing reperfusion is remarkable different for each of these thrombolytic agents, however the contribution in terms of reducing the overall mortality is equal for all the products. In addition reocclusion or prolonged 10 bleeding are frequent complications. This might be due to relatively low specificity for fibrin and the short plasma half-life of these compounds. Currently various application regimens and combinations of different fibrinolytic principles are tested in order to overcome some of the current short comings in thrombolytic therapie. The improvement which are expected are however rather small.

15 Recently a new group of patients occured with probelms such as acute thrombotic occlusion and late restenosis due to procedures such as angioplasty, atherectomy, arterial grafting or vessel wall stenting. The possible therapeutic interventions comprise anti-platelet, antithrombotic and thrombolytic strategies. Various other agents such as ticlopdine acting as ADP antagonists or Calcium 20 ionophore A-23187 and especially Asprin have a direct influence on the platelet function and have been suggested or used to prevent or minimize platelet aggregation. The new anti-platelet adhesion substance according to this invention could as well help to overcome these clinical complications when applied during surgery.

25 Another complication related to this topic arises if artificial surfaces come in contact with blood, then their is increased tendency to induce thrombotic events by activation of platelets and/or induction of coagulation. These effects may cause failure of vascular grafts, cardiac valves, stents, catheters or any other blood contacting device or material. The ability of the protein disclosed here to 30 create non-thrombogenic surfaces may therefore be further exploited by immobilization of this protein to the materials and devices described above. Such a treatment should render such materials or devices biocompatible and thromboresistant.

Due to the limitations associated with the available antithrombotic agents there is an actual need for new alternative strategies and therapeutics.

Background of the invention

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A potential for future improvements in the treatment of cardiovascular disorders may be contributed by approaches as disclosed in this invention which directly interfere with the collagen and/or vWF factor induced platelet adhesion.

Several novel inhibitors which prevent platelet adhesion are monoclonal
10 antibodies directed to vWF. It has as well been suggested that glycoprotein IIb/IIIa inhibitors may be beneficial in inhibiting platelet adhesion

Some of these inhibitors like the monoclonal Ab c7E3 have already been tested clinically while others like the KGD- and RGDF-inhibitors are still under study. However, the specificity of most of these new inhibitors is not very well studied ,
15 thus the spectrum of side effects that will be induced by using this inhibitors is still open and deserves careful examination.

A rich source for the screening of new compounds that interfere with collagen induced platelet adhesion is given in nature through blood-sucking animals. Several inhibitors have been isolated from nature as described in the literature: A
20 65 kD protein called Calin isolated from *Hirudo medicinalis* (US 5,587,360, WO 92/07005) (Munro, R., et al., Blood Coagulation and Fibrinolysis, 1991, 2, 179-184) and a 16kD (LAPP) protein isolated from the salivary glands of the leech *Haementeria officinalis* (US 5,324,715). Both proteins have been described as aggregation inhibitors as tested in static assays of collagen dependent platelet
25 aggregation.

Despite a proven in vitro activity LAPP failed to act in several well-established in vivo models (Schaffer L.W. et al.; Arterioscler. Thromb., 1993, 13, 1593-1601) and Connolly T.M. et al.; Thromb. Haemostas., 1993, 69, 589). The soft tick, *Ornithodoros moubata*, also contains an antiplatelet protein (Moubatin) which is
30 active in preventing collagen-stimulated platelet aggregation (Waxman, L. et al.; J. Biol. Chem., 1993, 268, 5445-49). Another inhibitor of platelet aggregation from a blood-sucking bug was disclosed in WO 9309137 by Noeske-Jungblut C. et al.. Smith et al. have isolated a 50 kDa protein from snake venom and a 19 kDa

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protein was isolated from a the saliva of *Triatoma pallidipennis*, a blood-sucking bug. The protein was found to contain a factor that specifically inhibits collagen-induced platelet aggregation. The 19 kDa protein named pallidipin inhibits collagen-mediated aggregation of platelets in plasma. No inhibition of aggregation
5 stimulated by other effectors (ADP, thrombin, thromboxane A2 mimetic U46619, phorbol ester) was detected. Pallidipin had no effect on platelet adhesion to collagen but inhibited ATP release from platelets. It interacted reversibly with platelets and may share with collagen a common target on them. The precise mechanism of action and therapeutic benefit of this protein is under investigation.
10 Gan et al. described Echistatin as an inhibitor binding to the fibrinogen receptor GP IIa/IIIb (J. Biol. Chem, 1988, 263, 19827-32).

Despite these exciting developments, the need continues to exist for supplying further anticoagulants and antithrombin which have increased efficacy in the inhibition of clot formation, vWF-induced platelet activation or endothelial cell
15 activation and which may be used pharmaceutically and produced in commercially feasible quantities.

Since none of the known proteins described sofar has developed into a compound with ideal therapeutic profile the inventors of the present invention decided to go ahead with a new screening strategy in order to detect more
20 relevant proteins.

Description of the Invention

25 In this invention an inhibitor isolated from *H. medicinalis* is described which directly acts one collagen-platelet interaction thus inhibiting platelet activation and early platelet-platelet interaction.

Sofare, there has not been a positive example in the literature which indicates that by using a screening approach that would exclude aggregation inhibitors as
30 well as lytic proteins from a source of naturally occuring compounds one could identify new anti-adhesive mechanisms or compounds. However this strategy was used in this invention. Since at least six different platelet surface

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glycoproteins are known to be involved in collagen adhesion and in addition several platelet derived compounds such as von Willebrand factor, fibronectin and thrombospondin have been shown to be involved as indirect mediators of collagen - platelet adhesion there has been little optimism in the beginning to
5 identify a new adhesion inhibitors.

Nevertheless, this approach was used to screen *Hirudo medicinalis* saliva knowing that not all the documented or unknown vWF related inhibitors as well as compounds directly acting on platelet receptors could be excluded. Thus the result of the screening was a surprise: A new protein named Saratin with anti-
10 adhesive activity for platelets which can be isolated from tissues and secretions of well investigated leech of the species *Hirudo medicinalis*.

The present invention comprises the active polypeptide Saratin isolated from the leech *Hirudo medicinalis*. The protein was isolated from saliva by a combination
15 of pressurized dialysis and at least one chromatographic step like anionic exchange chromatography and at least one reversed phase high performance chromatography (RP-HPLC) step. The pressure dialysis step turned out to be absolutely critical during the recovery of Saratin from saliva, since the strong concentration of saliva helped to overcome the otherwise tremendous loss of
20 bioactive Saratin. The isolated Saratin binds strongly to several collagens and blocks the adhesion of platelets to collagen coated such surfaces in a dose dependent fashion.

In order to optimize the screening cascade currently available techniques have
25 been developed to distinguish platelet adhesion and platelet aggregation: the ability of platelets to retard or stop flow through fibers, the contribution of platelets to in vitro clot formation, glass bead adhesion laboratories, or whole blood flows through the filter and platelet adhesion of anticoagulated platelet-rich plasma to filters composed of glass fibers or collagen under a regulated pressure gradient.

30

The protein (named Saratin) is characterized by the amino acid sequences depicted in sequence (SEQ. ID. NO. 1) and is constituted from 103 amino acids which make up a theoretical relative molecular weight of approximately 12068

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dalton \pm 1 kDa The protein exhibits a unique primary structure with no significant similarity to other previously described sequences. The protein is rich in aspartic and glutamic acid which contribute to the low isoelectric point of pH 3.7 ± 0.5 of the molecule as measured by IEF-PAGE.

5

SEQ. ID. NO. 1

E E R E D C W T F Y A N R K Y T D F D K S F K K S S D L D E C K K T C F K T
E Y C Y I V F E D T V N K E C Y Y N V V D G E E L D Q E K F V V D E N F T E
10 N Y L T D C E G K D A G N A A G T G D E S D E V D E D

- SDS-PAGE analysis demonstrated a strong shift in mobility upon reduction of the protein prior to electrophoresis, indicating posttranslational modifications.
- 15 Sequencing of the polypeptide had revealed six cysteine molecules which could make-up post-translational modifications of the protein. Electrospray mass spectrometry of Saratin revealed an actual molecular weight of 12061 indicating that up to three disulphide bonds are involved in the formation of the secondary structure of the native form of the protein.
- 20 The adhesion inhibitor according to the present invention is new because it differs from known aggregation inhibitors isolated from leeches especially from Calin or LAPP in the molecular weight, isoelectric point and amino acid sequence and biological activities.
- 25 The present invention provides as well isolated DNA comprising a polynucleotide encoding the leech derived platelet adhesion inhibitor having the amino acid sequence as shown for the protein. The nucleotide sequence representing the cDNA clone is shown in **SEQ. ID. NO. 2**, Position 1-63 of the nucleotide sequence represents a putative 21 amino acid leader sequence and position 64–
- 30 372 contains an open reading frame coding for a polypeptide of 103 amino acid residues and an amino acid sequence as shown for the mature protein in **SEQ. ID. NO. 1**.

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SEQ. ID. NO. 2

1

ATGAAGTATTTCTTGATTTCTTCCTTTGCCTCGCAAGCTTGCTGATCTCAAC

↓

5 TACTTCTTCAGAAGAACGTGAAGATTGTTGGACGTTTTACGCGAACAGAAAA

TATACAGACTTCGATAAATCTTTTAAGAAGTCCTCTGATCTTGACGAATGCAA

AAAAACATGTTTCAAGACGGAGTACTGCTACATCGTTTTTGAAGACACGGTC

10

AACAAGGAATGTTACTACAATGTCGTTGATGGTGAAGAGTTAGACCAAGAAA

AATTTGTTGTCGACGAAACTTCACGGAAAATTATTTGACAGACTGCGAGGG

15 TAAAGATGCAGGTAATGCGGCAGGTACAGGTGACGAGTCAGATGAAGTTGA

↓

TGAAGATTAA

The present invention also relates to recombinant vectors which include the
20 synthetic gene coding for the leech-derived platelet adhesion inhibitor of the
present invention, and a host cell containing the recombinant vectors. Methods
for recovering and isolating the expressed proteins have been based on tag-
technologies or have been adapted from the purification scheme developed for
the naturally occurring Saratin. Depending on the individual protocols used for
25 extracellular or intracellular expression in yeast cells, insect cells, baby hamster
kidney cells and E. coli cells transformed with the appropriate vectors the steps
for recovering the recombinant protein from the supernatant or sediments have to
be adapted by techniques known to the expert. Excellent expression was found in
E. coli as a host, where periplasmatic expression was contributed by insertion of
30 a pelB leader sequence. Products recovery from Escherichia coli (E. coli) was
achieved (around 5 mg/l) after osmolysis and centrifugation. Saccharomyces
cerevisiae (S. cerevisiae) (> 10 mg/l culture broth) with the alternative yeast
adopted vector was used in a paralleled experiment. The secreted material was

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isolated by centrifugation. Purification was achieved by cross-flow filtration and ion exchange chromatography. Expression was as will done in COS cells or CHO cells (750 ng/ml). The recombinant material proved to be pure and homogeneous by electrophoretic and chromatographic analysis and identical to saliva derived
5 Saratin as demonstrated by amino acid sequencing and molecular mass determination .

The invention also comprises methods of purifying the active protein from crude saliva of the leech and measuring its activity against platelets by static and dynamic assay methods as well as the use of this method to isolate rekombinant
10 protein.

Proteins of the present invention include variants which conserve the activity of the disclosed sequences, including fragments or subunits, naturally occurring variants, allelic variants, randomly generated artificial mutants and intentional sequence variations such as adding which conserve activity. Fragments or
15 subunits refers to to any portion of the sequence which contain fewer amino acids than the complete protein e.g. partial sequences excluding portions of the N- and/or C-termini of the complete protein.

The invention further covers hybrid proteins, such as fusion proteins or proteins resulting from the expression of multiple genes within the expression vector, and
20 may include a polypeptide having the specific activity of a disclosed protein linked by peptide bonds to a second polypeptide. Notably other variants of the proteins of the present invention are included, especially any variants that differ from the isolated protein only by conservative amino acid substitution. Such conservative amino acid substitutions are defined in Taylor et al., J. Mol. Biol., 1986, 188, 233.

25

Also included are methods for using the proteins to prevent or delay of platelet activation by inhibition of collagen-platelet interactions. The protein is useful in the prevention, prophylaxis, therapy and treatment of thrombotic diseases. Unlike all these previously described proteins, which act at various surface proteins on the
30 platelet, the protein from this invention has a unique mechanism of action. It binds tightly to the collagen surface and one mechanism of action is given by the coverage of specific collagen sides no longer available for interactions and binding of platelets. This type of new mechanism has the great advantage, that

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the platelets stay functionally intact during the application of the protein, so that very low or even no bleeding tendency is expected from this treatment.

Another important area of use is the treatment of various surfaces with the protein to render them non-adhesive for platelets and thereby create blood-compatible devices.

As indicated above, the polypeptides according to the present invention are suitable as pharmaceutically effective compounds in pharmaceutical compositions and combinations.

The pharmaceutical formulations according to the invention optionally may comprise additional active ingredients like Aspirin, anti-coagulants such as hirudin or heparin or thrombolytic agents such as plasminogen activator or streptokinase.

The novel polypeptide according to the invention may form pharmaceutically acceptable salts with any non-toxic, organic or inorganic acid. Inorganic acids are, for example, hydrochloric, hydrobromic, sulphuric or phosphoric acid and acid metal salts such as sodium monohydrogen orthophosphate and potassium hydrogen sulfate. Examples for organic acids are the mono, di and tri carboxylic acids such as acetic, glycolic, lactic, pyruvic, malonic, succinic, glutaric, fumaric, malic, tartaric, citric, ascorbic, maleic, hydroxymaleic, benzoic, hydroxybenzoic, phenylacetic, cinnamic, salicylic and sulfonic acids such as methane sulfonic acid. Salts of the carboxy terminal amino acid moiety include the non-toxic carboxylic acid salts formed with any suitable inorganic or organic bases. These salts include, for example, alkali metals such as sodium and potassium, alkaline earth metals such as calcium and magnesium, light metals of Group IIIA including aluminium, and organic primary, secondary and tertiary amines such as trialkylamines, including triethylamine, procaine, dibenzylamine, 1-ethenamine, N,N'-dibenzylethylene-diamine, dihydroabietylamine and N-alkylpiperidine.

30

As used herein, the term "pharmaceutically acceptable carrier" means an inert, non toxic solid or liquid filler, diluent or encapsulating material, not reacting adversely with the active compound or with the patient. Suitable, preferably liquid

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carriers are well known in the art such as steril water, saline, aqueous dextrose, sugar solutions, ethanol, glycols and oils, including those of petroleum, animal, vegetable, or synthetic origin, for example, peanut oil, soybean oil and mineral oil.

- 5 The formulations according to the invention may be administered as unit doses containing conventional non-toxic pharmaceutically acceptable carriers, diluents, adjuvants and vehicles which are typical for parenteral administration.

10 The term "parenteral" includes herein subcutaneous, intravenous, intra-articular and intratracheal injection and infusion techniques. Also other administrations such as oral administration and topical application are suitable. Parenteral compositions and combinations are most preferably administered intravenously either in a bolus form or as a constant fusion according to known procedures.

- 15 Tablets and capsules for oral administration contain conventional excipients such as binding agents, fillers, diluents, tableting agents, lubricants, disintegrants, and wetting agents. The tablets may be coated according to methods well known in the art.

20 Oral liquid preparations may be in the form of aqueous or oily suspensions, solutions, emulsions, syrups or elixirs, or may be presented as a dry product for reconstitution with water or another suitable vehicle before use. Such liquid preparations may contain conventional additives like suspending agents, emulsifying agents, non-aqueous vehicles and preservatives.

- 25 Topical applications may be in the form of aqueous or oily suspensions, solutions, emulsions, jellies or preferably emulsion ointments.

Unit doses according to the invention may contain daily required amounts of the protein according to the invention, or sub-multiples thereof to make up the
30 desired dose. The optimum therapeutically acceptable dosage and dose rate for a given patient (mammals, including humans) depends on a variety of factors, such as the activity of the specific active material employed, the age, body weight, general health, sex, diet, time and route of administration, rate of

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clearance, the object of the treatment, i. e., therapy or prophylaxis and the nature of the thrombotic disease to be treated, antiplatelet or anticoagulant activity.

Therefore, in compositions and combinations useful as antithrombotic in a treated patient (in vivo) a pharmaceutical effective daily dose of the peptides of this invention is between about 0.01 and 100 mg/kg body weight, preferably between 0.1 and 10 mg/kg body weight. According to the application form one single dose may contain between 0.5 and 10 mg of the thrombin inhibitor. To achieve an anticoagulant effect in extracorporeal blood a pharmaceutically effective amount of the inventive peptides is between 0.2 and 150 mg /l, preferably between 1mg and 20 mg/l extracorporeal blood.

It is also object of this invention to provide an implantable or extracorporeal medical device for use in contact with body fluids in order to render the device surface substantially thromboresistant, coated with an immobilized polypeptide as defined above and in the claims. The polypeptide according to the invention is immobilized on a medical device so as to render the surface biocompatible and thromboresistant. Such devices sometimes have surface properties which typically induce platelet aggregation, which is a disadvantage in their intended uses in implantable and extracorporeal devices in contact with blood or other body fluids. Examples for such devices which are commonly made from plastics materials and synthetic fibres are prostheses, artificial organs, ocular lenses, sutures, artificial vascular segments, catheters, dialysers, tubes and vessels carrying blood.

Posterior capsule opacification (PCO) is a common complication after cataract extraction, despite the modern surgical techniques and lenses which are used for this procedure. PCO is caused by the proliferation and migration of lens epithelial cells across the posterior capsule thus reducing the visual acuity. Physical treatments as well as chemically modified lenses have been proposed to reduce formation of PCO. Heparin lens coating or topical heparin eyedrops have been used to reduce PCO, indicating that thrombogenic mechanisms are involved in the formation of PCO.

Saratin has been shown to have significant advantages over heparin in preventing and blocking thrombogenicity. It is therefore another feature of this

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invention to provide a coating comprising Saratin which reduces thrombogenicity of the lens material used for refractive anterior or posterior chamber ocular implants which may be surgically implanted into the eye. This new type of coating avoids problems contributed by stimulated cell growth. In combination with other
5 medicaments which are for instance conferring cell death, Saratin coating helps to completely overcome posterior capsule opacification.

Brief Description of the Figures

Details of the figures are explained in examples 1 to 10.

Fig. 1

5 Separation of saliva components. Elution of Saratin is marked by *

a) shows the separation profile of saliva after DEAE anion exchange. Saratin fractions have been collected from peak 3 (Example 2).

10 b) shows re-chromatography of pooled fractions on Mono Q HR5/5. Samples have been collected from the last part of the major peak as indicates by bar (Example 2).

15 c) shows last chromatography step on semi-preparative analytical RP-HPLC of Saratin positive fractions collected from Mono Q HR5/5 (example 2). Active Saratin was recovered from the major peak (peak 3).

Fig. 2

20 SDS-PAGE of fractions collected from RP-HPLC. Saratin positive fraction is marked * (Examples 2 and 3).

Fig. 3

Linkersequence cloned in pASK75 (Example 6).

Fig. 4

25 E. coli expression vector for Saratin (Example 7).

Fig. 5

Baculo donor plasmid for Saratin (Example 7)

30

- 15 -

Fig. 6

Whole blood has been exposed to an artificial collagen surface and platelets adhesion has been visualized by staining. Saratin has been used as an inhibitor (Protein #607) Example 9

5

Fig. 7

Inhibition of platelet adhesion on collagen type III coated coverslips under shere conditions. Comparison of saliva and saratin. Example 9

10

Fig. 8

Saratin exhibits dose dependent inhibition of platelet binding adhesion to collagen type III coated coverslips under shere conditions. Example 9

15

Fig. 9

Yeast expression vector for Saratin (Example 13)

20

Detailed description of the invention**Example 1****25 Screening adhesion inhibitors**

Platelet adhesion to collagen has been the functional background four screening saliva components. In addition four additional tests available for the assesment of various actions of antithrombotic drugs such as the AZOCOLL assay, amidase
30 activity assay, vWillebrand dependent binding assay and platelet aggregation assay have been used to exclude functional properties which idealy would not be linked to an adhesion inhibitor. While most of this assays used here are standard assays, the platelet adhesion assay had to be modified to fitt our specific needs.

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In short: Horm collagen (Nycomed) has been coated to 96 well plates (Nunc) by using acidified collagen at a concentration of 20 µg /ml and incubating the plates overnight. After washing the plates 3 times with PBS the residual surface of the wells have been blocked with 1% BSA. Platelets isolated freshly from human
5 citrionylated blood have been added simultaneously with fractions derived from the individual column steps. Raw saliva, standardized to a protein concentration of xxx mg/ml, has been used as a standard to compare for inhibitory activity. The platelet inhibition assay turned out to be highly susceptible to buffer changes and elevated salt concentrations. Since all the samples have been processed by ion-
10 exchanger chromatography the direct testing of the fractions in the inhibition assay turned out to be complicated and unreliable. Therefore all samples to be tested have been applied to a Centricon based concentration step which simultaneously reduced ionic strenght and supports concentration prior to measurement.

15

Example 2

Purification of a natural inhibitor

The invention used saliva collected from *H. medicinalis* which is known to contain
20 a number of bioactive proteins such as hirudin, elastase inhibitors, collagenases and platelet aggregation inhibitors such as calin (Munro, R. et al.; Blood Coagulation and Fibrinolysis, 1991, 2, 179-184) and LAPP (Schaffer, L.W. et al.; Arterioscler. Thromb., 1993, 13, 1593-1601). Besides this characterized proteins the majority of the roughly eighty proteins detectable by SDS-PAGE are still
25 unknown. In the present invention the fractionated saliva and the screening strategy as described in Example 1 was used in order to screen for the novel proteins that interferes directly with platelet collagen interaction.

Separation of an adhesion inhibitor activity from raw saliva turned out to be a
30 critical task, mainly due to the irreversible loss of most of the adhesion inhibitory activity in the first chromatographic step. Additives such as 12% ethanol and divalent cations as recommended by Munro, R. et al did not improve the situation. However the high salt concentration of the saliva in combination with the low

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overall protein concentration (2 mg/ml) asked for an initial concentration or buffer exchange step. Thus several strategies for enrichment, concentration or buffer exchange were explored. Traditional dialysis led to the complete loss of activity. Most of the other standard techniques such as ion exchangers, affinity columns, size exclusion (loss was irrespective of the column resin) failed independent of the nature of the separation technology or buffer and additives used. Surprisingly it turned out, that pressurized dialysis of 500ml saliva was a successful method to concentrate the saliva proteins (about 30-40 times) and simultaneously get rid of the unwanted buffer components. Unexpectedly the saliva raw material processed in this way was an ideal starting material for further purification and the recovery of bioactivity anti-adhesive components out of the saliva was no longer a real problem. Since cation exchangers or affinity columns turned out to be an insufficient step in purification, weak anion exchanger such as DEAE-Fastflow or EMD-DEAE-Fractogel were used. 12% ethanol and divalent cations have been tested, however the results were similar with or without this additives. Further optimization of the chromatographic steps led to a sequence of DEAE-column, CM- column and as a final step an reversed phase RP18 column. Optimization of the chromatographic conditions have been performed on a Biacore chromatographic system using analytical columns available from Pharmacia. The gradients used to operate the DEAE-column, CM- column and RP18 columns are given in Figures 1a, b, c. The Biacore based separation has been scaled up by using FPLC techniques. The optimized running conditions have been directly converted into the a semi-preparative scale of separation by following the instruction of the manufacturer with the exception of the RP18 column step which was maintained with Biacore technique in order to minimize loss of purified material. Recovery of purified protein from the last chromatographic RP-step was done by speed vacuum centrifugation. Samples collected from the last RP-step have been collected and analysed by SDS-PAGE (Figure 2) Subsequently samples have been resuspended in PBS and used to perform analytical as well as functional tests. Typically a yield of about 750 µg/l Saratin was recovered from unprocessed saliva.

Example 3

Biochemical characterization

5

Purification of Saratin as disclosed in Example 1 led to an essentially pure protein with an apparent molecular weight around 21 kDa shown under reducing conditions in SDS-PAGE (Figure 2). The complete amino acid sequence was obtained by direct sequencing of the first 48 amino acids of the purified protein and was completed by the sequencing of several internal peptides generated via enzymatic degradation. The complete protein sequence is disclosed in **SEQ. ID. NO. 1**.

The protein is composed of 103 amino acids with an calculated molecular weight of 12067,9 and an actual molecular weight of 12061,9 as deduced by ESI-Mass spectrometry. This difference in theoretical and measured molecular weight indicates that all the six cysteines identified in Saratin are involved in the formation of S-S bridges. This finding is supported by the observation of a strong change in mobility of the protein when chromatographed in SDS-PAGE under reducing or non-reducing conditions. Furthermore the protein is rich in acidic acids such as Glu and Asp. Isoelectric focussing using IEF-PAGE (Immobiline) technique revealed an isoelectric point of $\text{pH } 3.7 \pm 0.5$. In a comparative study we have used the purified leech protein as a reference and compared it with the physicochemical properties of recombinant Saratin derived from baculo, yeast and E.coli expression. All three proteins turned out to be identical in their properties.

Characterization of protein by SDS-PAGE visualized by Coomassie (Figure 2) or silver-staining and/or Western blot analysis revealed, that the protein was homogeneous and in a non-glycosylated form.

30 Example 4

mRNA Preparation and cDNA Synthesis

- 19 -

RNA was prepared from the medicinal leech, *Hirudo medicinalis*; using guanidinium thiocyanate method. mRNA was purified from total RNA using "Oligotex mRNA kit" (QIAGEN).

- 5 cDNA was synthesized using "Marathon cDNA Amplification kit" (CLONTECH). The DNA sequence encoding *Saratin* was initially amplified using PCR oligonucleotide primers as follows:

5'-TTCTAGAATTCAGCGGCCGC(T)₃₀N₁N-3' (N₁=G,AorC)

10 EcoRI Not I

After cDNA synthesis, a universal adaptor was ligated to both ends of the cDNA. The sequence of the universal adaptor is as follows:

5'-CCATCCTAATACGACTCACTATAGGGC-3' Primer AP1

15

5'-ACTCACTATAGGGCTCGAGCGGCC-3' Primer AP2

5'-TAATACGACTCACTATAGGGCTCGAGCGGCCCGCCCGGGCAGGT-3

T7 Promoter

3'-H₂NCCCGTCCA-PO₄-5'

20

Example 5

Amplification and isolation of the *Saratin* gene by PCR

25

Several degenerated primers were synthesised based on the *Saratin* amino acid sequence. 3'-RACE PCR and 5'-RACE PCR were carried out using one Neo-Calix primer and one universal primer AP1 or AP2. PCR products were cloned in TA cloning vector pCR2.1 or pCR Script SK(+) vector and sequenced. After
30 sequencing both the 3'-RACE PCR fragment and 5'-RACE PCR fragment, the *Sarastatin* full length gene sequence was obtained. Amplifying the Neo-Calix gene by PCR using the primers from both the 3'-end and the 5'-end of the Neo-Calix gene, a full length of *Saratin* gene was obtained.

- 20 -

DNA Sequencing was performed with more than 15 different PCR clones. However only one significant change which caused amino acid sequence change was found in only one clone. It is very likely that this change was caused by PCR. Another five silent changes which do not cause amino acid sequence change were found. Thus it is unlikely that these changes were caused by PCR, because the same changes were found in different clones.

The Saratin gene ORF is 372bp and contains a 21 amino acid signal sequence and 103 amino acid sequence coding for the mature protein. The amino acid sequence deduced from the PCR clone was found to be identical to the sequence obtained by sequencing of the natural saliva derived protein.

Example 6

15 Expression in COS cells and the detection of expressed protein

In order to express Saratin gene in mammalian cells such as COS or CHO, Saratin gene was cut from the vector pCR Script SK(+) using XhoI+XbaI and cloned into mammalian cell expression vector pCI-neo (Promega). pCI-neo was chosen because it contains T7 and T3 promoter for in vitro expression and neomycin resistant gene for G418 selection, and can be used in both COS and CHO cells.

In addition to the signal sequence and mature protein sequence, the insert contains a Koz. sequence at the 5'-end for efficient translation and a MRGS(H)₆ tag. at the 3'-end (C-Ter-minal) for detection of protein expression, purification and concentration. The plasmid constructure was named pNC-31.

pNC-31 plasmid DNA was used for the transfection of COS cells. The COS cells growing at log phase were washed twice with PBS and dissolved in PBS at a concentration of 1×10^7 /ml. Then 12ug plasmid DNA (less than 50µl in H₂O or TE buffer) was added in 0.7-0.8 ml COS cells suspension and mixed in a electroporation cuvette. Electroporation was performed at 1.9k-v, 25uFD for 10 min and transfered to a 90mm plate. After adding 8 ml DMEM medium containing

- 21 -

10% FCS and antibiotics the cells were grown for three days. The supernatantes and the cells have been used for further isolation of protein and detection. The expressed protein was detected by western blotting method using anti-MRGS(H)6 antibody. Purification was performed using chelators such as NTA or
5 imido acetic acid immobilized on a column matrix and modified with metall ions such as Co, Ni, or Cu.

Example 7

10 Construction of the *E. coli* expression vector and expression

Because of the variable codon usage in different biological systems, some codons are used very rarely in *E. coli*. In order to enable expression an optimized version in *E. coli* the gene had to be converted to *E. coli* codon usage according
15 to standard procedures.

The expression in *E. coli* was performed using a modified version of the plasmid pASK75, which carries the tet promoter region. (Skerra, A., et al. Gene, 1994, 151, 131 - 135). The modification was made by cloning a new linker between the
20 XbaI an Hind III sites (Figure 3). The new linker contains the ompA leader sequence, another multiple cloning site and a 6xHis-tag instead of the strep-tag.

To construct the expression vector for Saratin it was necessary to introduce 5' Cla I and 3' Eco47III restrictionsites by PCR method.

25 Therefore the two primers 5' GCATCGATGGAAGAACGTGAAGAC and 3' TAGCGCTTTTGACGTCGTCGTCA were used. The PCR product first was cloned into the PCR II vector system (Invitrogen) and sequenced.

In a second step the Saratin gene was cloned into the modified pASK75 vector using the rectrictionsites 5'ClaI and 3' HindIII.

30 After expressing and proving the activity of this recombinant Sarastatin in a second PCR reaction the His-tag was removed and the start codon of the Sarastatin gene was directly fused to the omp A leader sequence. The primers

- 22 -

for this PCR reaction were: 5' GAAGAATGCAAGGATGAGGATTATTG and
3' AAGCTTCTAGTCTTCGTCAACTTCG.

As an example for the expression in *E. coli* the expression vector pRG72 (Figure
5 4), which contains the structural gene of Saratin fused to the ompA leader
sequence, was transformed into W3110 competent cells. Cells have been
induced after they have been grown to mid-log phase. 1 hour thereafter the
recombinant Sarastatin could be clearly detected.

10

Example 8

Construction of the Baculo Donor Plasmid and expression

15 For expression of Saratin in the Baculo virus expression system the Bac-To-
Bac™ Baculovirus Expression System from Gibco Life Technologies was used.
To get a selection system the Honeybee melitin leader sequence was fused to
the Saratin gene and to introduce the restriction sites 5' BamHI and 3' KpnI one
single PCR reaction was carried out using the 5' Primer: CGG ATC CAT GAA
20 ATT CTT AGT CAA CGT TGC CCT TGT TTT TAT GGT CGT ATA CAT TTC
TTA CAT CTA TGC GGA AGA ACG TGA AGA TTG TTG GAC T
and the 3' Primer: GGT ACC TCA CAT ATC TTC ATC AAC

The corresponding PCR product was cloned into the PCR II Vector (Invitrogen)
and sequenced. Then the Melitin - Saratin fusion was cloned into the pFastBac
25 vector using the restriction sites 5' BamHI and 3' KpnI.

Generation of recombinant baculoviruses and Saratin expression was performed
with the Bac-To-Bac Expression System. The donor plasmid pTD13 was
transformed into DH10Bac competent cells which contain the bacmid with a mini-
attTn7 target site and the helper plasmid. The mini-Tn7 element on the donor
30 plasmid can transpose to the a mini-attTn7 target site on the bacmid in the
presence of transposition proteins provided by the helper plasmid. Colonies
containing recombinant bacmids were identified by disruption of the *lacZ* gene.
High molecular weight mini-prep DNA is prepared from selected *E. coli* clones

- 23 -

containing the recombinant bacmid, and this DNA was then used to transfect insect cells. Detailed descriptions are included in the instruction manual of the expression kit.

5

Example 9

Platelet Adhesion to Collagen Under Flow Conditions (Dynamic assay)

- 10 In the platelet adhesion assay, whole human blood is perfused through a parallel flow chamber to examine the adhesive activity of platelets to a collagen coated coverslips under high shear flow (simulating in vivo arterial conditions) as originally described by Sakariassen et al (Meth. Enz., 1988, 169, 37-70). Human placental collagen, type III (Sigma), solubilized in 50 mmol/L acetic acid was
15 sprayed onto cleaned glass coverslips (18mm x 18mm) with a retouching airbrush. Coverslips were stored in PBS at 4°C overnight.

- Fresh whole human blood (anticoagulated with low molecular weight heparin; 20 U/ml) was prewarmed at 37°C for 10 min before being used. Preparations of proteins according to this invention were pipetted onto coverslips (30 µl per
20 coverslip) and incubated for 10 min in a humid chamber at room temperature before being inserted into the perfusion chamber. The blood was allowed to perfuse through the chamber (at 37°C for 5 min at a shear rate of 1300 s⁻¹.)

- Subsequently, the coverslips are removed, washed in PBS and fixed in 0.25%
25 glutaraldehyde for 30 min., and thereafter stained with May-Grünwald Giemsa. Figure 9 shows a typical expamle. Extensive coverage with stained platelets is seen in the untreated control surface. A comparable surface pretreated with Saratin shows dramatically deminished (by 80%) platelet binding. Platelet adhesion was quantified with a light microscope (magnification x 1000) coupled to
30 a computerized image analyzer (Leica). Results were expressed as the percentage of the surface covered with platelets and platelet aggregates.

Comparison of saliva inhibitory activity has been compared to purified protein as demonstrated in Figure 10. Platelet adhesion on collagen type III coated

- 24 -

coverslips at a shear rate of 1300 s⁻¹ with crude saliva (#616) produced an inhibition of about 48% compared to control. Purified protein (#607; Saratin) demonstrated a reduction in platelet deposition of about 81% at standardized protein concentrations. The inhibition increases in a dose-dependent manner with increasing concentrations of purified protein (Figure 11).

Example 10

Immunization and antibodies

10

With the first lot of highly purified natural protein available, immunization of animals has been started right away. Immune sera were raised in rabbits and high titered reagents were available for further screening. Additional antisera became available when the peptide sequence of the complete protein was available and three synthetic peptides have been synthesized (amino-acid sequence 83-103, 13-30, 58-69) coupled to KLH and used for immunization. Three sera directed to an N-terminal peptide segment and two sera specific for C-terminal peptides have established. With the high titered immune sera available it has been possible to establish and use Elisa technology to monitor and quantitate purification of naturally purified as well as recombinant protein. Thus the time consuming and rather work intensive platelet-inhibition assay could be replaced and was applied only to confirm the inhibitory potential of finally purified protein.

25 Example 11

Immuno-Assays for estimation of Saratin binding

Acidified Horm-collagen (Nycomed) has been used for the coating of 96-well micro-titer plates (Nunc) 50 µl of the collagen solution (20 µg/ml) has been used to coat plates over night. Prior to testing the plates have been washed three times with PBS and have been incubated with a BSA solution (1%) in order to prevent non-specific adhesion. 50 µl Saratin has been added in serial delution and have

- 25 -

been incubated for one hour. Plates were washed three times prior to the application of anti-Saratin antibody for detection. After an additional one our incubation step surplus antibody has been removed and a second biotin-labeled Antibody has been used for detection. Read-out has been performed via
5 streptavidin-POD catalyzed colour reaction with substrates such as ODB-tablets (Dako) measured at 490 nm.

Example 12

10

Competition assay for screening inhibitors

Recombinant tagged Saratin (His-tag) prepared as described in Example 7 has been compared with native untagged Saratin for collagen binding. Plates coated
15 with acidified Horm-collagen (Nycomed) have been prepared as described in Example 11. Detection was performed with rabbit anti-Saratin antibodies. The tagged and untagged Saratins showed identical binding properties. Alternatively the untagged Saratin version has been modified by biotinylation (Pierce, biotinylation kit) and compared to unmodified Saratin. Binding properties to
20 collagen have been identical. Furthermore experiments have been performed using the biotinylated

Saratin to cross-compete with unmodified Saratin, peptides, saliva derived Saratin, complete saliva or antibodies directed to Saratin. Binding of the various „competitors“ to collagen has been tested by estimating the binding of biotinylated
25 Saratin using a streptavidine-POD conjugate and ODB-substrate reaction for detection. This assay comprising biotinylated Saratin was typically used for the estimation of Saratin concentration in saliva (750µg/l), epitope mapping of Saratin directed antibodies, evaluation of bioactive Saratin, mutated Saratin. In order to explore the potential of the assay blocking as well as none blocking anti-Saratin
30 antibodies raised with specific Saratin peptides have been used.

Example 13**Yeast expression vector and expression**

The pichia multi copy expression system (Invitrogen) has been used as a typical
5 example for yeast expression. The construction of the yeast expression vector is
shown in Fig. 9. For generation of the expression vector for Saratin the PCR
amplification method has been used to generate restriction ends (5' EcoR I,
3' Not I) compatible with the ligation into the appropriate vector (pPIC9K). The the
following primers have been used:

10 5' GCA TGC GGC CGC CTA ATC TTC ATC AAC TTC
 3' GCA TGA ATT CGA AGA ACG TGA AGA TTG

Before transforming the *Pichia* spheroplasts the expression vector has been
linearized with Sal I. Colonies have been screened for His⁺ Mut⁺-positiv mutants
to ensure integration of the Saratin gene. Typical growing conditions have been:
15 28–30°C, up to an optical density of OD 2-6. Induction of expression was
performed after resuspending centrifuged cells in medium and addition of
methanol up to a final concentration of 0,5% and maintaining this condition for an
extended time periode which typically would be 24 hours. After a typical
fermentation periode of 6 days the product is recovered from the supernatant and
20 analysed by SDS-PAGE and ELISA.

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Patent Claims

1. A polypeptide isolated from *H. medicinalis* having a molecular weight of about 12 000 \pm 1kD with the biological activity of an inhibitor of collagen-dependent
5 platelet adhesion.
2. A polypeptide of claim 1 having an isoelectric point of pH 3.7 \pm 0.5
3. A polypeptide of claim 1 or 2 having six cysteine molecules capable of forming
10 -S-S- bridges
4. A polypeptide of claim 1 which comprises an amino acid sequences of SEQ. ID. NO. 1
- 15 5. A polypeptide comprising an amino acid sequence according to claim 4 which is at least 80% identical to the amino acid sequence of SEQ. ID. NO. 1 over its entire length.
6. An isolated polynucleotide encoding a polypeptide of claims 1-5.
20
7. An isolated polynucleotide comprising a DNA sequence of SEQ. ID. NO. 2, or a DNA sequence complementary to said DNA sequence wherein said polynucleotide is encoding a polypeptide of claim 1-5.
- 25 8. The polynucleotide of claim 7 wherein said polynucleotide comprises a DNA sequence that is at least 80% identical to that of SEQ ID NO: 1 over its entire length.
9. An expression vector comprising a DNA sequence of claims 6-8.
- 30 10. A host cell comprising the expression vector of claim 9.
11. An expression system comprising a host cell of claim 10.

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12. A process for producing a polypeptide of claims 1-5 comprising a host of claim 10 culturing said host under conditions sufficient for the production of said polypeptide and recovering the polypeptide from the culture supernatant or cell residue.

5

13. An antibody immunospecific for a polypeptide of claims 1-5.

14. A pharmaceutical formulation which comprises a polypeptide according to claims 1-5 and a pharmaceutical acceptable carrier or excipient therefore.

10

15. A pharmaceutical active agent of claim 14 for the treatment of thromboembolic processes

16. A pharmaceutical formulation of claims 14 or 15 comprising additional drugs wherein the additional drug is selected from aspirin, heparin or streptokinase or a combination thereof.

17. Use of a polypeptide according to claims 1-5 for the manufacture of a medicament for the treatment of thromboembolic diseases.

20

18. Use of a polypeptide according to claims 1-5 for coating artificial surfaces.

19. Use of a polypeptide according to claims 1-5 for modifying intraocular lenses in order to lessen the thrombogenicity of the lens material.

25

20. Use of a polypeptide according to claims 1-5 for contacting the lens surface

21. Use of a polypeptide according to claims 1-5 for covalent crosslinking to modify said lens material.

30

22. Use of antibodies according to claim 13 and polypeptide according to claim 1-5 to measure samples derived from claim 12 or a treated subject.

- 29 -

23. A method for identifying compounds which inhibit (antagonize) or agonize the polypeptide of claim 1 -5 by observing the binding, or stimulation or inhibition of a functional response.

5 24. An agonist identified by the method of claim 23.

25. An antagonist identified by the method of claim 23.



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Fig. 1a

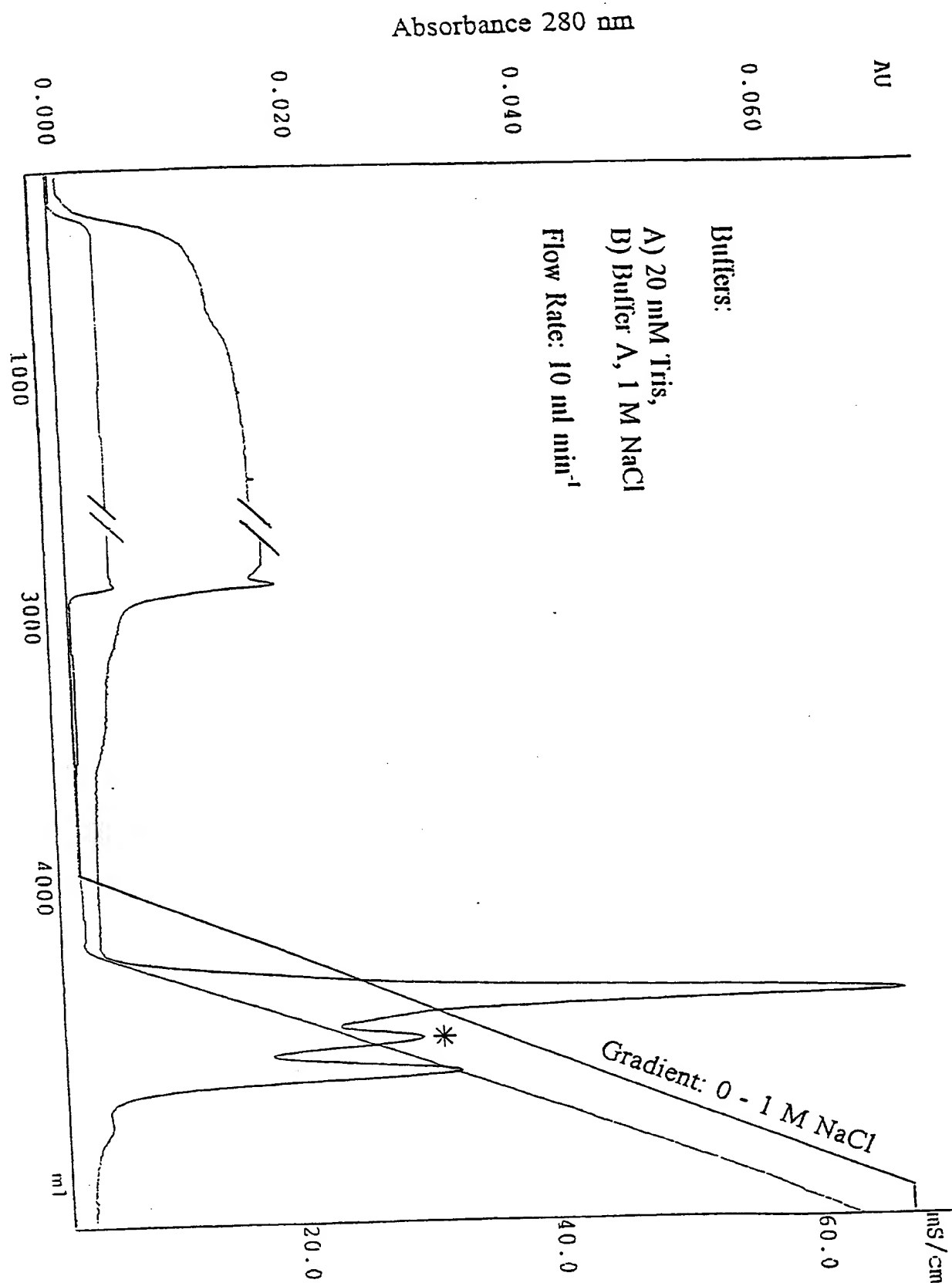


Fig. 1b

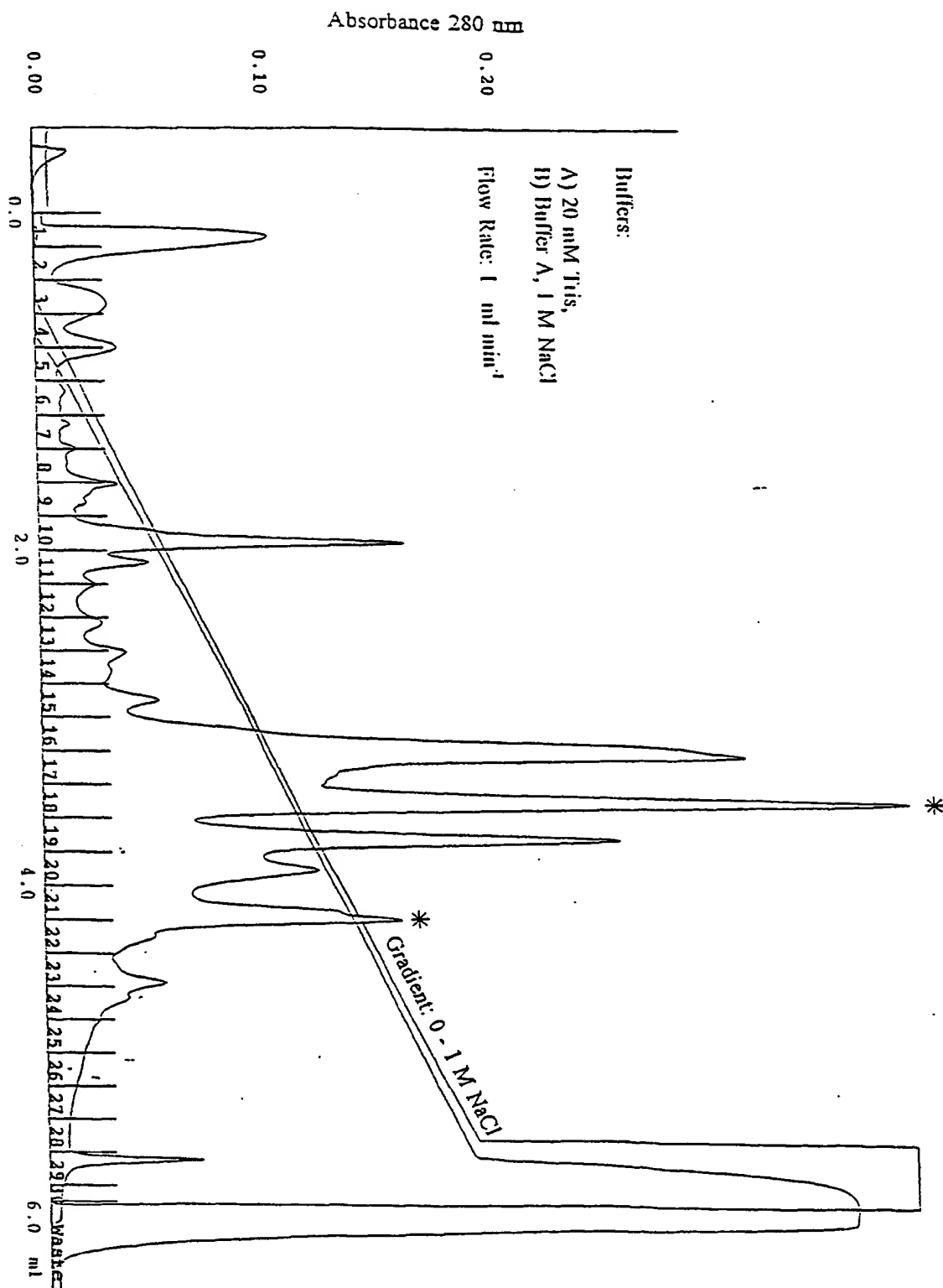


Fig. 1c

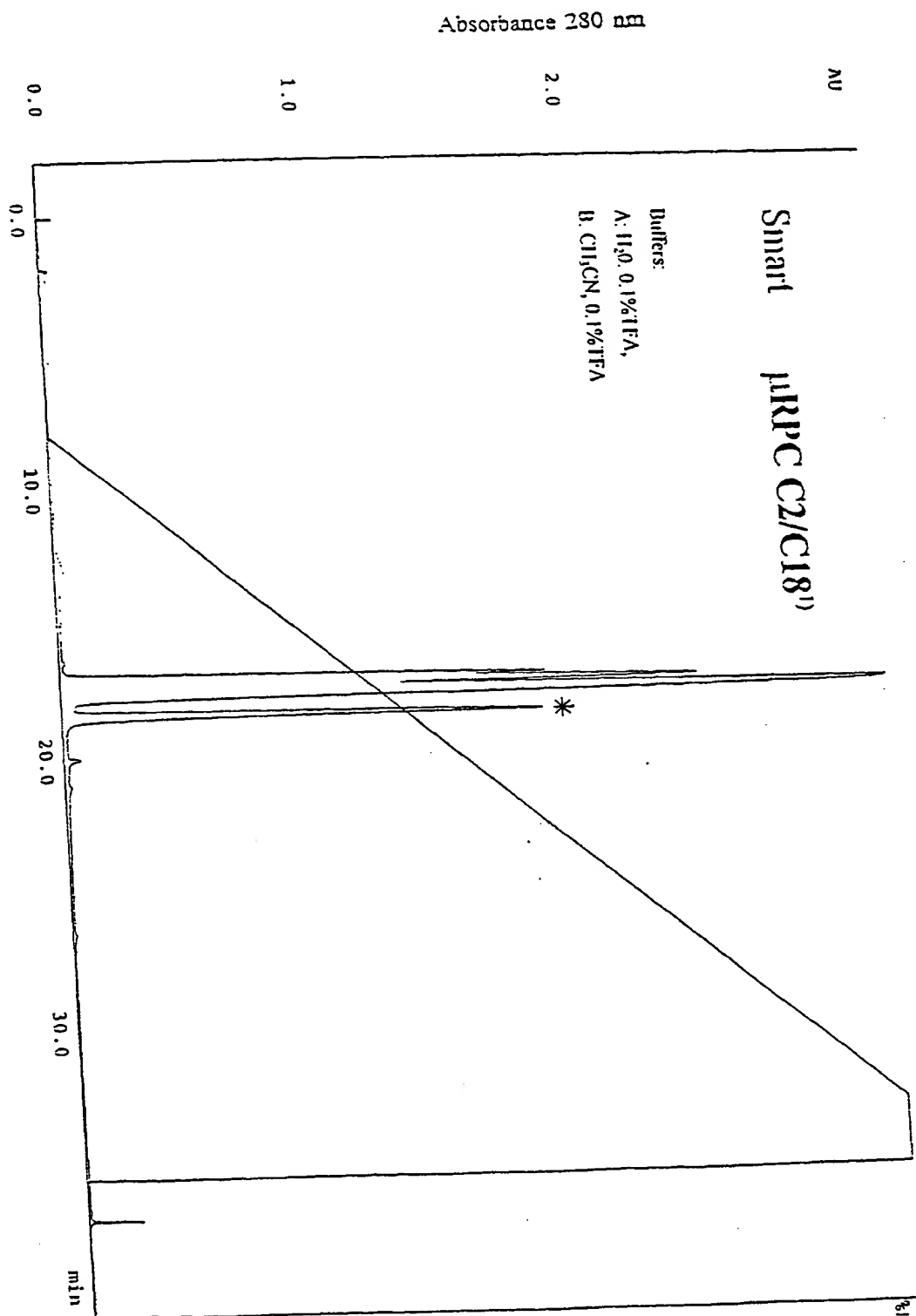


Fig.2

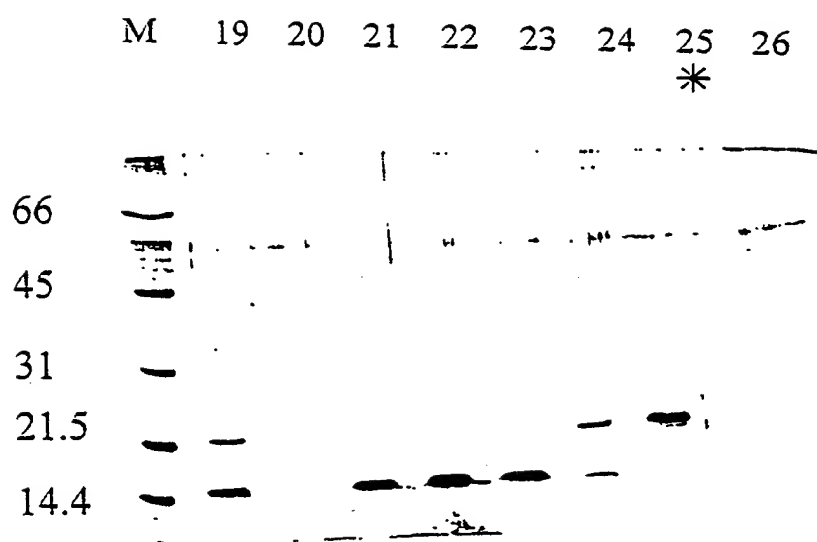


Fig. 3

Xba I

119 CTAGATAACG AGGGCAAAAA ATGAAAAAGA CAGCTATCGC GATTGCAGTG GCACTGGCTG
 TATTGC TCCGGTTTTT TACTTTTTCT GTCGATAGCG CTAACGTAC CCGTACCGAC
 1 MetLysLysT hrAlaIleAlaIleAlaValAlaLeuAlaG
 ClaI EcoRI Sall KpnI SmaI BamHI

179 GTTTCGCTAC CGTAGCGCAG GC AT CGA TGA ATT CGA GCT CGG TAC CCG GCG
 CAAAGCGATG GCATCGCGTC CG TA GCT ACT TAA GCT CGA GCG ATG GCG CCC
 14 IyPheAlaThrValAlaGlnAla
 XhoI Sall PstI Eco47III

230 ATC CCT CGA GGT CGA CCT GCA GCG AGC GGTATGAGAGSATCCCATCACCATCACCA
 TAG GGA GGT CCA GGT GGA CGT CCG TCG CGATAGTCTCCTAGCGTAGTGCTAGTGGT
 Hind III 1 AlaMetArgGlySerHisHisHisHisHis

286 TCACTAATAGA
 AGTGATTATCTTCGA
 10 sHis.....

Fig.4

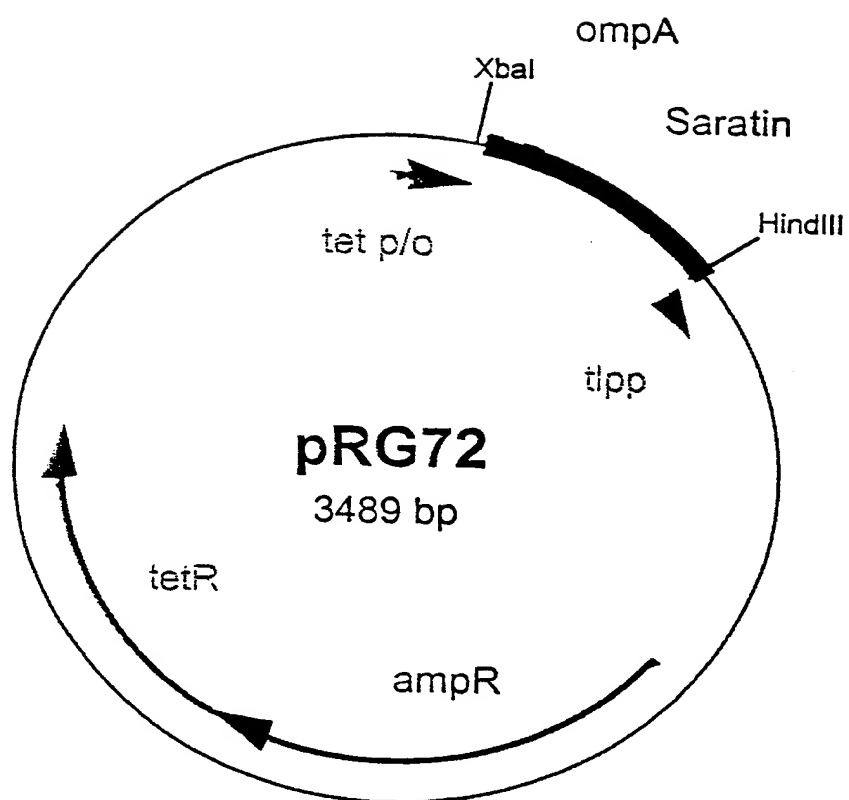


Fig.5

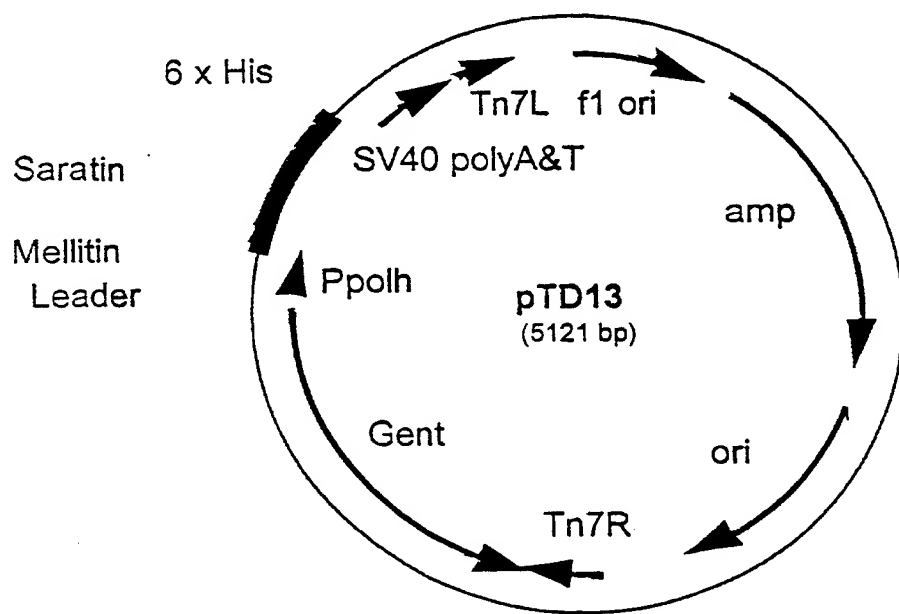


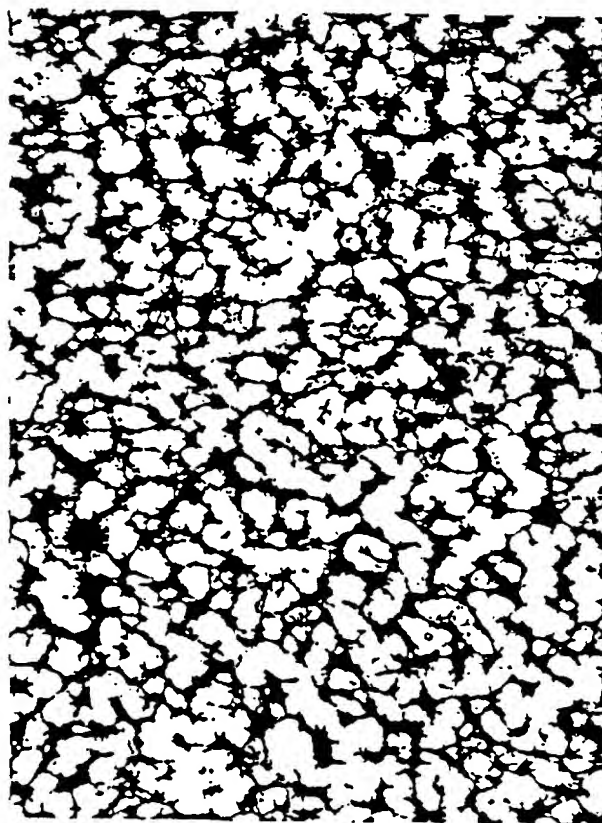
Fig.6

Protein # 607



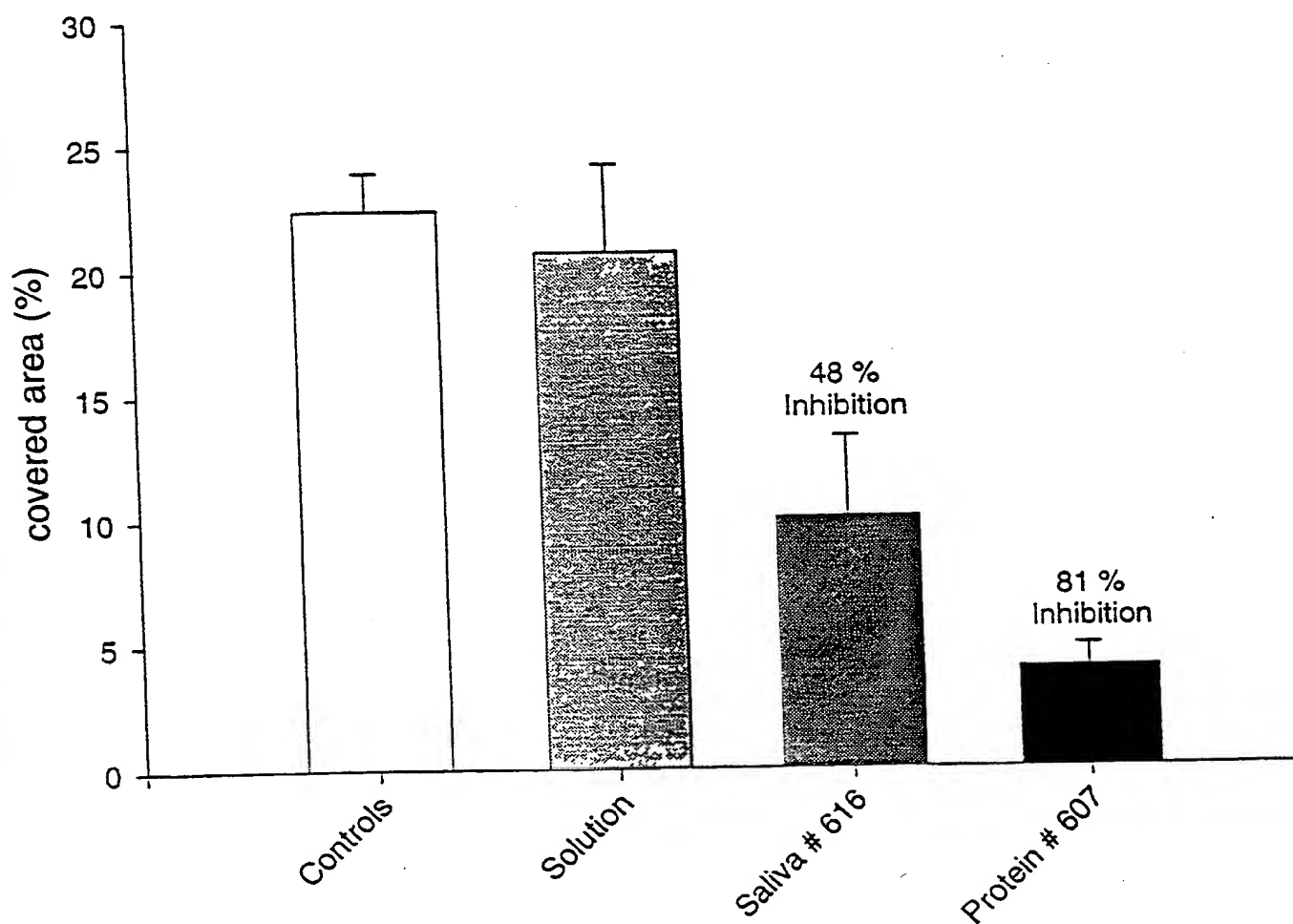
80 % Inhibition

Control



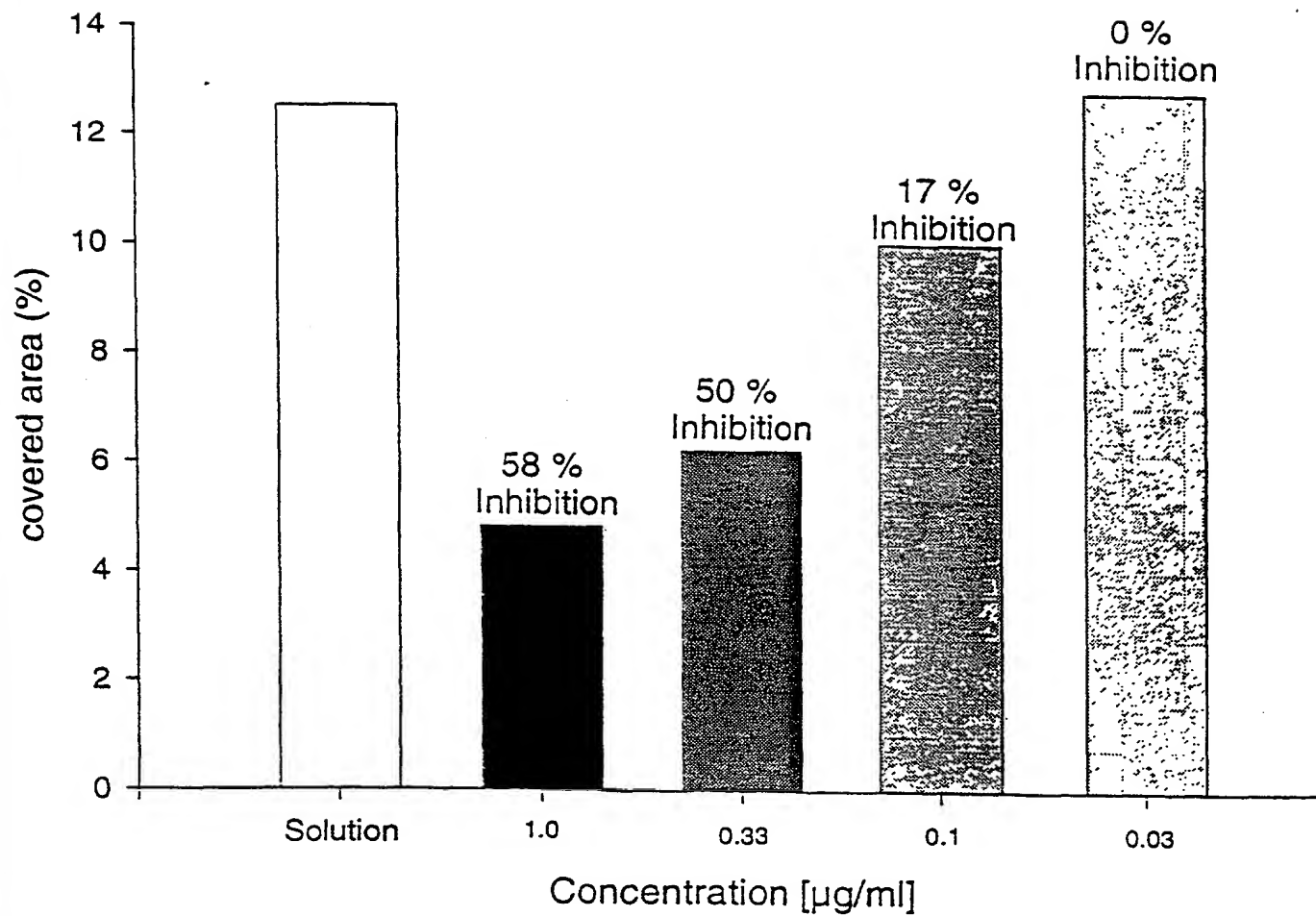
Shear rate: 1300 s^{-1}
Human Blood
Collagen Type III from Human Placenta
30 μl /coverslip

Fig.7



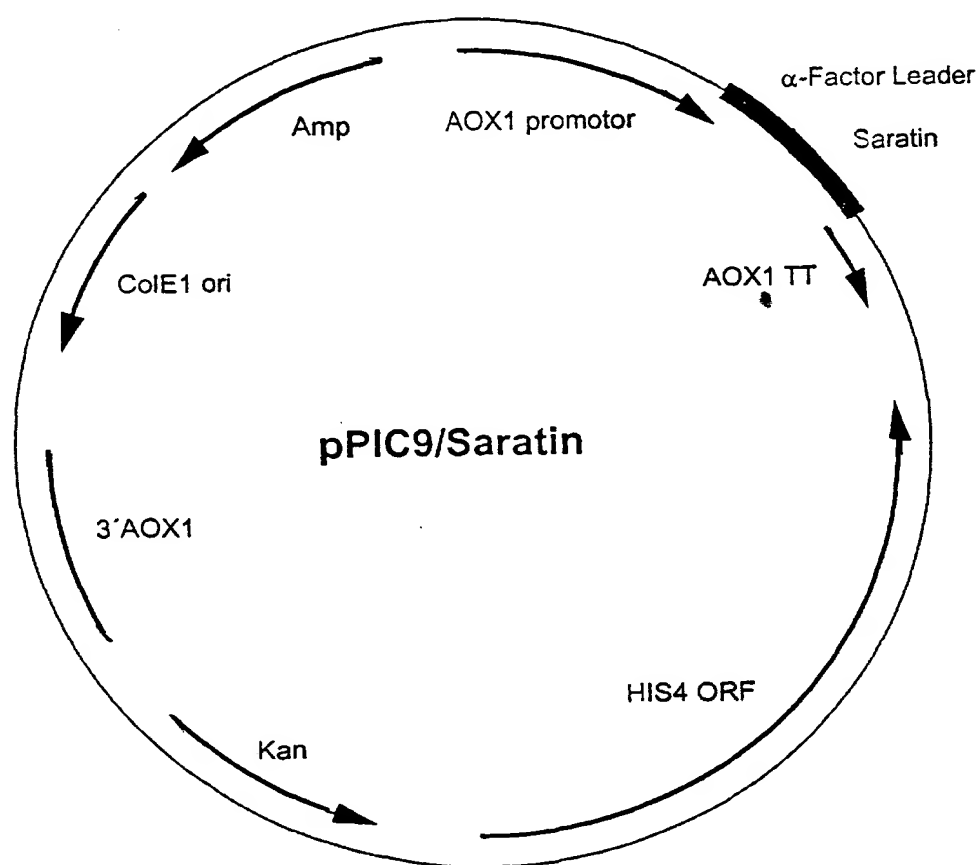
Collagen Type III from Human Placenta
Saliva # 616 0.11 mg/ml: 30 μ l on coeverslip n=4
Protein # 607 0.11 mg/ml: 30 μ l on coverslip n=2

Fig.8



Protein # 607
Collagen Type III from Human Placenta

Fig. 9



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Abstract:

A naturally occurring protein isolated from the saliva of the medicinal leech *Hirudo medicinalis* is described which strongly binds to collagen thus acting as an inhibitor of natural platelet adhesion to collagen. The protein has a molecular weight of about 12 000 Dalton, an acidic isoelectric point and contains six cysteines. The protein was sequenced and the gene was cloned from a *H. medicinalis* cDNA-library. Procedures for producing such polypeptide by recombinant techniques are disclosed. The recombinant and the natural occurring proteins are potent inhibitors of collagen-dependent platelet adhesion and therefore useful for the therapeutic treatment of various conditions related to hard disease and diseases of the circulation system. Furthermore the protein is useful for coating natural or artificial collagen surfaces in order to render them nonadhesive for cells and prevent the activation of cells.

